

**K830529 BIOFEEDBACK-TRAINING INSTRUMENT**Mar 24, 1983  
34 days to decisionK830529 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k830529/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Feb 18, 1983
Decision date	Mar 24, 1983
Days to decision	34 days
Third-party review	No

**APPLICANT**

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Company	<b>Relaxomat, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1995

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Device record: <https://www.510kdatabase.net/k830529/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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